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(54) **Liquid copolymers of epsilon-caprolactone and lactide**

Flüssige Copolymere aus E-Caprolacton und Lactide

Copolymères liquides à partir d'E-caprolactone et de lactide

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Description

FIELD OF THE INVENTION

[0001] This invention relates to absorbable liquid copolymers of ϵ -caprolactone and lactide. More specifically, it relates to liquid copolymers of caprolactone and lactide suitable for use in biomedical applications such as drug delivery, coatings for surgical sutures and needles and lubricants for medical devices.

BACKGROUND OF THE INVENTION

[0002] Pharmaceuticals have been conventionally administered, either in oral or parenteral dosage forms. However, these dosage forms are not always well suited for particular drugs or prolonged drug therapies. Many pharmaceuticals cannot be administered orally and patient noncompliance with dosage instructions is also a significant problem with oral dosage forms. To overcome these and other shortcomings of these dosage forms several new dosage forms have recently been developed.

[0003] The most notable recent development has been the use of bioerodible or bioabsorbable polymer matrices as carriers for implantable or intrauterine devices. Several publications describe these materials such as U.S. Patent 4,304,767 to Heller et al., "Biodegradable block copolymer matrices for long-acting contraceptives with constant release" J. Contr. Rel. 32 (1992) 3-14 by Z.W. Gu et al., and U.S. Patent 5,030,457 to Ng.

[0004] Heller describes a family bioabsorbable poly(ortho esters) which may be used as a matrix for controlled drug release. The polymers Heller et al. describes are synthesized by reacting a polyol with a diketene acetal. The polymers produced by this synthesis tend to be rigid because of the pentaerythritol segments in the polymer backbone. Unfortunately, this limits the use of these polymers to solid implants which generally must be surgically implanted.

[0005] Similarly, Gu et al. describes hard microspheres triblock copolymer of poly(ϵ -caprolactone-co-DL-lactide-co-glycolide) for the controlled release of contraceptives. The microspheres that Gu et al. describes are designed to be injected thereby avoiding the need to surgically implant a solid dosage. Unfortunately, these copolymers must be formed into microspheres and the kinetics of the pharmaceutical release are complicated by the different release mechanisms of the individual blocks of the triblock copolymer.

[0006] Ng describes a bioerodible polymer poly(ortho ester) formed from a one step reaction of an ortho ester and a triol. These polymers are more flexible than the polymers described by Heller and can be employed in ointments, gels and creams. Unfortunately the poly(ortho esters) described by Ng are highly susceptible to acid hydrolysis which limits their utility as a carrier for acidic

pharmaceuticals.

[0007] EP-A-509203 discloses copolymers prepared by reaction of lactide to a second monomer e.g. caprolactone, valerolactone, decalactone or hydroxybutyric acid whereby the molar ratio of lactide is from 95 to 70 and of the second monomer is from 5 to 30 and that dependent on the molar ratio of the monomers the copolymers are from highly viscous to solid compounds.

[0008] Thus it would be a significant contribution to the art to provide a bioabsorbable polymer that is easy to administer and slowly hydrolyses as it releases a drug.

SUMMARY OF THE INVENTION

[0009] In one aspect, the invention is a pharmaceutical carrier comprising a random copolymer of from 55 to 70 percent by mole lactide and from 45 to 30 mole percent ϵ -caprolactone. These liquid copolymers may be used as carriers for a variety of pharmaceutical compounds.

[0010] The copolymers of this invention are also useful for other biomedical applications. For example, the copolymers may be used as coatings for surgical sutures, surgical needles or as lubricants for medical devices such as trocars.

DETAILED DESCRIPTION OF THE INVENTION

[0011] The random copolymers of this invention are composed of lactide and ϵ -caprolactone within a range suitable to provide a liquid at room temperature. The amount of lactide contained in the copolymer carriers of this invention will be in the range of from about 55 to about 70 mole percent and the amount of ϵ -caprolactone will be in the range of from about 45 to 30 mole percent of the copolymer (where the total of the mole percents is 100). Preferably, the amount of lactide in the copolymer will be in the range of from 55 to 65 mole percent and the amount of ϵ -caprolactone will be in the range of from 45 to 35 mole percent (where the total of the mole percents is 100).

[0012] The copolymer carriers of this invention are typically characterized by being liquids at room temperature (25°C). The copolymers of the present invention have an intrinsic viscosity as determined in a 0.1 g/dl solution of hexafluoroisopropanol (HFIP) at 25°C ranging from about 0.05 to less than 0.8, preferably from about 0.05 to about 0.3 and most preferably from 0.05 to 0.2. A copolymer with an intrinsic viscosity below 0.05 may fail to significantly impart a controlled release profile to a pharmaceutical, and a carrier copolymer with an intrinsic viscosity above 0.8 may be too viscous to be easily administered.

[0013] The copolymers of this invention may be mixed with one or more therapeutic agents. The preferred dosage forms for the copolymer of the invention are sustained release parenterals, bioerodible ointments, gels,

creams, and similar soft dosage forms adapted for the parenteral or topical administration of therapeutic agents, other modes of administration (e.g., transdermal) and compositional forms (e.g., more rigid transdermal forms) are within the scope of the invention as well.

[0014] Parenteral administration of a bioerodible composition of the invention can be effected by either subcutaneous, or intramuscular injection. Parenteral formulations of the copolymer may be formulated by mixing one or more pharmaceuticals with a liquid copolymer. Other suitable parenteral additives may be formulated with the copolymer and pharmaceutical active, however, if water is to be used it should be added immediately before administration. The bioerodible ointment, gel or cream may also be injected as is or in combination with one or more suitable auxiliary components as described below. Parenteral delivery is preferred for administration of proteinaceous drugs such as growth factors, or growth hormone.

[0015] The bioerodible ointments, gels and creams of the invention will include: an ointment, gel or cream base comprising one or more of the copolymers described herein and a selected therapeutic agent. The therapeutic agent, whether present as a liquid, a finely divided solid, or any other physical form, is dispersed in the ointment, gel or cream base. Typically, but optionally, the compositions include one or more other components, e.g., nontoxic auxiliary substances such as colorants, diluents, odorants, carriers, excipients, or stabilizers.

[0016] The amount of active agent will be dependent upon the particular drug employed and condition being treated. Typically the amount of drug represents about 0.001% to about 70%, more typically about 0.001% to about 50%, most typically about 0.001% to about 20% by weight of the total composition being common.

[0017] The quantity and type of copolymers incorporated into the parenteral, ointment, gel, cream, is variable. For a more viscous composition, a higher molecular weight polymer is used. If a less viscous composition is desired, a lower molecular weight polymer can be employed. The product may contain blends of the liquid or low melting point copolymers to provide the desired release profile or consistency to a given formulation.

[0018] While not essential for topical or transdermal administration of many drugs, it may in some cases, with some drugs, be preferred that a skin permeation enhancer be coadministered therewith. Any number of the many skin permeation enhancers known in the art may be used. Examples of suitable enhancers include dimethylsulfoxide (DMSO), dimethylformamide (DMF), N,N-dimethylacetamide (DMA), deslymethylsulfoxide (C¹⁰MS)), ethanol, eucalyptol, lecithin, and the 1-N-dodecylcyclazacycloheptan-2-ones (available under the trademark Azone ® from the Nelson Research and Development Company, Irvine, California).

[0019] The variety of different therapeutic agents which can be used in conjunction with the copolymers of the invention is vast. In general, therapeutic agents

which may be administered via the pharmaceutical compositions of the invention include: antiinfectives such as antibiotics and antiviral agents; analgesics and analgesic combinations; anorexics; antihelmintics; antiarthritics; antiasthmatic agents; anticonvulsants; antidepressants; antidiuretic agents; antidiarrheals; antihistamines; antiinflammatory agents; antimigraine preparations; antinauseants; antineoplastics; antiparkinsonism drugs; antipruritics; antipsychotics; antipyretics, antispasmodics; anticholinergics; sympathomimetics; xanthine derivatives; cardiovascular preparations including calcium channel blockers and beta-blockers such as pindolol and antiarrhythmics; antihypertensives; diuretics; vasodilators including general coronary, peripheral and cerebral; central nervous system stimulants; cough and cold preparations, including decongestants; hormones such as estradiol and other steroids, including corticosteroids; hypnotics; immunosuppressives; muscle relaxants; parasympatholytics; psychostimulants; sedatives; and tranquilizers. Suitable pharmaceuticals for parenteral administration are well known as is exemplified by the Handbook on Injectable Drugs, 6th edition, by Lawrence A. Trissel, American Society of Hospital Pharmacists, Bethesda, Maryland, 1990.

[0020] In two particularly preferred embodiments the therapeutic agents for administration in conjunction with the bioerodible polymers of the invention are antibacterial agents for the treatment of deep wounds, and antibiotics for periodontal treatment (e.g., tetracycline). Other preferred drugs for use with the presently disclosed polymers include proteinaceous drugs such as epidermal growth factors or growth hormones.

[0021] Depending on dosage form, the pharmaceutical compositions of the preceding section may be administered in different ways, i.e., parenterally, or topically. Preferred dosage forms are liquid dosage forms which can be administered parenterally.

[0022] The copolymers, upon contact with body fluids including perspiration, or saliva (depending upon the mode of administration), undergoes gradual bioerosion with concomitant gradual exposure of the dispersed drug to the afflicted tissue. This can result in prolonged delivery (over, say 1 to 10,000 hours, preferably 2 to 1000 hours) of effective amounts (say, 0.0001 mg/kg/hour to 10 mg/kg/hour) of the drug. This dosage form can be administered as is necessary depending on the subject being treated, the severity of the affliction, and the judgment of the prescribing physician.

[0023] Topical application can be enhanced by occlusion, i.e., placing a barrier over the area treated so as to enhance absorption into the skin. Topical administration is preferred for wound healing and in the treatment of periodontal disease.

[0024] The following examples illustrate the claimed invention.

Example 1**LIQUID COPOLYMERS OF ϵ -CAPROLACTONE/L(-) LACTIDE @ 45/55 BY MOLE PERCENT INITIAL COMPOSITION**

[0025] A flame dried, 250 ml, round bottom single neck flask was charged with 51.36 gm (0.45 mole) of ϵ -caprolactone, 79.27 gm. (0.55 mole) of L(-)lactide, 0.735 milliliters of propylene glycol (USP grade), and 0.101 milliliters of stannous octoate (0.33 molar in toluene). The flask was fitted with a flame dried mechanical stirrer. The reactor flask was purged with nitrogen three times before venting with nitrogen. The reaction mixture was heated to 160°C and maintained at this temperature for about 20 hours. The copolymer was dried under vacuum 13,33 Pa (0.1 mm Hg) at 110°C for about 7 hours to remove any unreacted monomer. The copolymer had an inherent viscosity of 0.22 dl/g in hexafluoroisopropanol (HFIP) at 25°C. The copolymer was a liquid at room temperature. The mole ratio of poly[ϵ -caprolactone]/poly[lactide]/caprolactone/propylene glycol ester was found to be 44.8/50.8/1.2/3.2 by NMR.

Example 2**LIQUID COPOLYMERS OF ϵ -CAPROLACTONE/L(-) LACTIDE @ 40/60 BY MOLE PERCENT INITIAL COMPOSITION**

[0026] The procedure in Example 1 was substantially repeated, except that 45.66 gm (0.40 mole) of ϵ -caprolactone, 86.48 g (0.60 mole) of L(-) lactide were used. The copolymer was dried under vacuum 13,33 Pa (0.1 mm Hg) at 110°C for about 7 hours to remove any unreacted monomer. The copolymer had an inherent viscosity of 0.38 dl/g in hexafluoroisopropanol (HFIP) at 25°C. The copolymer was a liquid at room temperature. The mole ratio of poly[ϵ -caprolactone]/poly [lactide]/caprolactone/propyleneglycol ester was found to be 40.0/54.2/1.9/3.9 by NMR.

Example 3**LIQUID COPOLYMERS OF ϵ -CAPROLACTONE/L(-) LACTIDE @ 35/65 BY MOLE PERCENT INITIAL COMPOSITION**

[0027] The procedure in Example 1 was substantially repeated, except that 39.95 gm (0.35 mole) of ϵ -caprolactone, 93.68 g (0.65 mole) of L(-) lactide are used. The copolymer was dried under vacuum 13,33 Pa (0.1 mm Hg) at 110°C for about 7 hours to remove any unreacted monomer. The copolymer had an inherent viscosity of 0.19 dl/g in hexafluoroisopropanol (HFIP) at 25°C. The copolymer was a liquid at room temperature. The mole ratio of poly[ϵ -caprolactone]/poly[lactide]/caprolactone/propyleneglycol ester was found to be 35.6/54.2/1.2/3.7

by NMR.

Example 4**LIQUID COPOLYMERS OF ϵ -CAPROLACTONE/L(-) LACTIDE @ 45/55 BY MOLE PERCENT INITIAL COMPOSITION**

[0028] The procedure in Example 1 was substantially repeated, except that 6.0 milliliters of glycerol (USP grade) was used instead of 0.735 milliliters of propylene glycol. The copolymer was dried under vacuum 13,33 Pa (0.1 mm Hg) at 110°C for about 7 hours to remove any unreacted monomer. The copolymer had an inherent viscosity of 0.12 dl/g in hexafluoroisopropanol (HFIP) at 25°C. The copolymer was a liquid at room temperature. The mole ratio of poly[ϵ -caprolactone]/poly [lactide]/caprolactone/glycerol ester was found to be 46.5/43.6/2.2/7.7 by NMR.

Example 5**LIQUID COPOLYMERS OF ϵ -CAPROLACTONE/L(-) LACTIDE @ 40/60 BY MOLE PERCENT OF INITIAL COMPOSITION**

[0029] The procedure in Example 4 was substantially repeated, except that 45.66 gm (0.40 mole) of ϵ -caprolactone, 86.48 g (0.60 mole) of L(-) lactide were used. The copolymer was dried under vacuum 13,33 Pa (0.1 mm Hg) at 110°C for about 7 hours to remove any unreacted monomer. The copolymer had an inherent viscosity of 0.11 dl/g in hexafluoroisopropanol (HFIP) at 25°C. The copolymer was a liquid at room temperature. The mole ratio of poly[ϵ -caprolactone] /poly[lactide]/caprolactone/glycerol ester was found to be 40.4/49.0/1.7/8.9 by NMR.

Example 6**LIQUID COPOLYMERS OF ϵ -CAPROLACTONE/L(-) LACTIDE @ 35/65 BY MOLE PERCENT INITIAL COMPOSITION**

[0030] The procedure in Example 4 was substantially repeated, except that 39.95 gm (0.35 mole) of ϵ -caprolactone, 93.68 g (0.65 mole) of L(-) lactide were used. The copolymer was dried under vacuum 13,33 Pa (0.1 mm Hg) at 110°C for about 7 hours to remove any unreacted monomer. The copolymer had an inherent viscosity of 0.12 dl/g in hexafluoroisopropanol (HFIP) at 25°C. The copolymer was a liquid at room temperature. The mole ratio of poly[ϵ -caprolactone] /poly[lactide]/caprolactone/glycerol ester was found to be 36.3/54.5/1.3/7.9 by NMR.

Claims

1. A liquid copolymer comprising from 55 to 70 mole percent lactide and from 45 to 30 mole percent ϵ -caprolactone, wherein the liquid copolymer is in liquid form at 25°C and has an intrinsic viscosity between 0.05 dl/g and less than 0.8 dl/g as determined in a 0.1 g/dl solution of hexafluoroisopropanol at 25°C.
2. The copolymer of claim 1 wherein the amount of lactide in the copolymer is in the range of 55 to 65 mole percent and the amount of ϵ -caprolactone is in the range of 45 to 35 mole percent.
3. The copolymer of claim 1 or claim 2 wherein the intrinsic viscosity of the copolymer is between 0.05 dl/g and 0.3 dl/g.
4. Pharmaceutical composition comprising the copolymer of any one of claims 1 to 3 admixed with one or more therapeutic agents.
5. Pharmaceutical composition according to claim 4 in a dosage form selected from sustained release parenterals, bioerodible ointments, gels and creams.
6. A surgical suture coated with a copolymer according to any one of claims 1 to 3.
7. A surgical needle coated with a copolymer according to any one of claims 1 to 3.
8. A medical device lubricated with a copolymer according to any one of claims 1 to 3.

Patentansprüche

1. Flüssiges Copolymer, umfassend von 55 bis 70 Molprozent Lactid und von 45 bis 30 Molprozent ϵ -Caprolacton, dadurch gekennzeichnet, daß das flüssige Copolymer bei 25°C in flüssiger Form vorliegt und eine intrinsische Viskosität zwischen 0,05 dl/g und weniger als 0,8 dl/g aufweist, wie bestimmt in einer 0,1 g/dl Lösung von Hexafluoroisopropanol bei 25°C.
2. Copolymer nach Anspruch 1, dadurch gekennzeichnet, daß die Menge an Lactid in dem Copolymer in dem Bereich von 55 bis 65 Molprozent und die Menge an ϵ -Caprolacton in dem Bereich von 45 bis 35 Molprozent liegt.
3. Copolymer nach Anspruch 1 oder 2, dadurch gekennzeichnet, daß die intrinsische Viskosität des Copolymers zwischen 0,05 dl/g und 0,3 dl/g liegt.

4. Pharmazeutische Zusammensetzung, umfassend das Copolymer nach einem der Ansprüche 1 bis 3, beigemischt mit einem oder mehreren therapeutischen Agenz(ien).
5. Pharmazeutische Zusammensetzung nach Anspruch 4 in einer Dosierungsform ausgewählt aus Parenteralien mit verzögerter Freigabe, bioerodierbaren Salben, Gels und Cremes.
6. Chirurgische Naht, beschichtet mit einem Copolymer nach einem der Ansprüche 1 bis 3.
7. Chirurgische Nadel, beschichtet mit einem Copolymer nach einem der Ansprüche 1 bis 3.
8. Medizinische Vorrichtung, geschmiert mit einem Copolymer nach einem der Ansprüche 1 bis 3.

Revendications

1. Un copolymère liquide comprenant de 55 à 70 moles pour cent de lactide et de 45 à 30 moles pour cent de ϵ -caprolactone, où le copolymère liquide est sous forme liquide à 25°C et présente une viscosité intrinsèque entre 0,05 dl/g et moins de 0,8 dl/g tel que déterminée dans une solution à 0,1 g/dl d'hexafluoroisopropanol à 25°C.
2. Le copolymère de la revendication 1, dans lequel la quantité de lactide dans le copolymère est dans la gamme de 55 à 65 moles pour cent et la quantité de ϵ -caprolactone est dans la gamme de 45 à 35 moles pour cent.
3. Le copolymère de la revendication 1 ou de la revendication 2, dans lequel la viscosité intrinsèque du copolymère est entre 0,05 dl/g et 0,3 dl/g.
4. Composition pharmaceutique comprenant le copolymère d'une quelconque des revendications 1 à 3 mélangé avec un ou plusieurs agents thérapeutiques.
5. Composition pharmaceutique selon la revendication 4 dans une forme galénique choisie parmi produits parentéraux à libération prolongée, pommades bioérodables, gels et crèmes.
6. Une suture chirurgicale revêtue d'un copolymère selon une quelconque des revendications 1 à 3.
7. Une aiguille chirurgicale revêtue d'un copolymère selon une quelconque des revendications 1 à 3.
8. Un dispositif médical lubrifié avec un copolymère selon une quelconque des revendications 1 à 3.